AD-A280 333 ENTATION PAGE

Form Approved

OMB No. 0704-0188

mated to average 1 neur per response, including the time for reviewing instructions, searching dust sources, or reviewing the collection of internation. Send comments regarding this purcen estimate or any other aspect of sing in purcen, to washington reasoursers Services, Directorate for information Operations and Reports, 1215 Jetterson in Office of Management and Sudget, Paperwork Reduction Project (07C4-2181), Washington, DC 20563.

II ACCUCY HEE DAILY MADERALL IS DESCRIPTION TO THE PROPERTY OF								
	1. AGENCY USE ONLY (Leave bia		2. REPORT DATE	1	3. REPORT TYPE AND DATES COVERED Journal article			
- 1-	TITLE AND SUSTITUE					INDING NUMBERS		
	Low risk of sexual transm	3. 70	MOING HOMBERS					
- 1			•		Pr	_62787A		
1					,	 2 - 3M162787A870		
6	. AUTHOR(S)					A - AR		
	Watts DM, Corwin AL, Omar	MA. H	vams KC			T: _1288		
	, , , , , , , , , , , , , , , , , , , ,	,	, uu	410		•		
				171C				
7	PERFORMING ORGANIZATION Naval Medical Research		8. PERFORMING ORGANIZATION REPORT NUMBER					
1	Commanding Officer							
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) Naval Medical Research Institute Commanding Officer 8901 Wisconsin Avenue Retherda Manufacial 20000 5000						NMRI 94-18		
Bethesda, Maryland 20889-5607								
1_				14	(
9.	SPONSORING / MONITORING A	GENCY	NAME(S) AND ADL	(ES)	10. SP	ONSORING / MONITOR	ING	
1	Naval Medical Research a	d	AGENCY REPORT NUMBER					
	National Naval Medical C	Center			1			
Building 1, Tower 12						DN243536		
8901 Wisconsin Avenue								
	Bethesda, Maryland 20889	-5606						
11	SUPPLEMENTARY NOTES						······································	
1	Reprinted from: Transact	ions o	f the Royal Society	of Tropical Medi	cine and Hy	viene 1994 Valor		
	pp.55-56	5	,	-	crite dild riye	1334 VO1. 88	•	
_								
12:	a. DISTRIBUTION / AVAILABILITY	STATE	MENT		126. DI	STRIBUTION CODE		
١,	Approved for public relate							
Approved for public release; distribution is unlimited.								
					Accesion For			
					NTIS C	RA&I	7	
13	ABSTRACT (Meximum 200 word	e/c l			- DTIC T	7 *		
		-3/			Unannou	nced 🗀		
						ication		

					Ву	ibution /		
					Distribution			
					Ava	ilability Codes	1	
					·		_	
					Dist '	Ivail and/or Special	- 1	
						Opecial I	1	
1	_10/70						ł	
	-18470				A-1 3	27)		
				•		-01		
				16	1 /	145		
	グリ		<i>"</i>	ŧU.	14	140		
14.	SUBJECT TERMS	•				15. NUMBER OF FA	GES -	
sexually transmitted diseases, viral hepatitis, hepatitis C, epidemiology,								
Sé	erology	ea, VI	ioi nepatitis, nepa	titis C, epidemic	ology,	16. PRICE CODE		
		٠.						
	SECURITY CLASSIFICATION 1		URITY CLASSIFICATION	19. SECURITY CLA		20. LIMITATION OF	ESTRACT	
	OF REPORT nclassified		THIS PAGE	OF ABSTRACT	•			
O,	ירימסטדובת	Un	classified	Unclassified		Unlimited		

16

Low risk of sexual transmission of hepatitis C virus in Somalia

Douglas M. Watts¹, Andrew L. Corwin², Mahmoud A. Omar³ and Kenneth C. Hyams¹ ¹Naval Medical Research Institute, Bethesda, Maryland, USA; ²US Naval Medical Research Unit No. 3, Cairo, Egypt; ³Ministry of Health, Mogadishu, Somalia

Abstract

The prevalence in Somalia of antibody to hepatitis C virus (anti-HCV) was determined in a survey of 236 female prostitutes, 80 sexually transmitted disease (STD) clinic patients, 79 male soldiers, and 43 tuberculosis patients. Of 98 (22%) serum samples repeatedly anti-HCV reactive by first and second generation enzymelinked immunosorbent assay kits, only 8 (1-8%) were anti-HCV positive by immunoblot assay (RIBA-2). Anti-HCV seropositivity by immunoblot assay was not associated with any risk group or with positive syphilis serology (found in 18% of subjects) or antibody to human immunodeficiency virus 1 (in 1-4% of subjects). These data indicate that sexual transmission of hepatitis C virus is not common in Somalia among sexually active populations, including female prostitutes and other groups at high risk of STDs and the acquired immune deficiency syndrome.

Introduction

Hepatitis C virus (HCV) has been found in developed countries to be a common cause of post-transfusion hepatitis and a common infection among groups with frequent parenteral exposure, such as illicit drug abusers (ALTER et al., 1989a; ESTEBAN et al., 1990; VAN DEN HOEK et al., 1990). The role of sexual transmission in the spread of HCV is not as well understood, particularly in developing countries, although HCV does not seem to be transmitted by sexual contact as easily as hepatitis B virus (ALTER et al., 1989b; EVERHART et al., 1990).

In developing countries, HCV infection has not been found as frequently as hepatitis B infection; however, high-risk groups in these countries have not been comprehensively evaluated (AL-FALEH et al., 1991; JACKSON et al., 1991; SAEED et al., 1991). In this investigation, populations living in Somalia and at high risk of sexually transmitted diseases (STDs) and human immunodeficiency virus (HIV) infection were investigated for HCV infection.

Patients and Methods

During 1990, the Somali Ministry of Health surveyed 438 subjects living in 3 major urban centres in Somalia: Mogadishu, Merca, and Chismayu. Subjects included 236 female prostitutes, 80 STD patients (69% male), 79 male military personnel, and 43 patients (58% male) with Mycobacterium tuberculosis infection. The mean age of the entire population was 28 years (range 13-79 years). All subjects were chosen sequentially during clinic or hospital visits on the days that the study was conducted; no selection criterion was used.

A serum sample was obtained from all study subjects and tested for total anti-HCV antibody using both first and second generation commercial enzyme-linked immunosorbent assay (ELISA) kits (Abbott Laboratories, Abbott Park, Illinois, USA). Repeatedly reactive sera were further verified with a second-generation immunoblot assay (RIBA-2[®]; Chiron Corporation, Emeryville, California, USA). Sera reactive by both ELISA and RIBA-2 were considered to be positive for anti-HCV and to represent active infection (FOLLETT et al., 1991). Sera were also screened for HIV-1 antibody by ELISA, and repeatedly reactive samples were confirmed by Western blotting. Lastly, sera were tested for syphilis infection by the rapid plasma reagin card test, with confirmation by the fluorescent treponemal antibody-absorption test (FTA-ABS).

Results

Among 438 study subjects, 83 (18.9%) had serum samples repeatedly reactive for anti-HCV by first-generation ELISA. By second generation ELISA, 74 (16.9%) sera were repeatedly reactive for anti-HCV: 59 serum samples Address for correspondence: Dr. K. C. Hyams, Naval Medical

Address for correspondence: Dr K. C. Hyams, Naval Medical Research Institute, 12300 Washington Avenue, Rockville, MD 20852, USA.

which were also reactive by first-generation assay and 15 additional sera which were negative by first-generation ELISA. Among all 98 repeatedly ELISA-reactive sera, only 8 (1.8%) were anti-HCV positive and 6 (1.4%) were indeterminate by RIBA-2. All 8 RIBA-positive sera were identified by the second generation ELISA; however, 4 of the 8 were missed by the first-generation ELISA. Sera from 79 subjects (18.0%) were positive by FTA-ABS and 6 (1.4%) were positive for HIV-1 antibody by Western blotting.

There was no significant association between HCV infection, determined by RIBA-2, and the age, sex, or risk group of study subjects (Table). The mean age of the 8

Table. Prevalence of antibodies to hepatitis C virus and human immunodeficiency virus 1, and positive syphilis serology, among 436 subjects living in Somalia and at high risk of sexually transmitted diseases and ADS

Risk group	No. examined	Number positive (percentages in parentheses) Anti-HCV ^a Anti-HIV-1 ^b FTA-ABS ^c			
Female prostitutes	236	4 (1.7)	5 (2·1)	73 (30.9)	
STD clinic patients	80	2 (2.5)	0 (-)	3 (3.8)	
Military personnel	79	1 (1.3)	1 (1.3)	3 (3.8)	
Tuberculosis patients	43	1 (2.3)	0 (-)	0 (-)	

^{*}Immunoblot for anti-hepatitis C antibody.

subjects positive for anti-HCV was 32 years, compared to a mean age of 28 years for the other study subjects. None of the anti-HCV positive subjects had HIV-1 antibody, and only 3 of 8 subjects with anti-HCV were FTA-ABS positive.

Discussion

Based on currently available tests for anti-HCV, these data indicate that sexual transmission of HCV is not common in Somalia among sexually active populations, including female prostitutes and individuals with anti-HIV-1 antibody and positive syphilis serology. A low risk of HCV infection has been found in other prostitute populations in developing countries, suggesting that HCV is not readily transmitted by sexual contact even among highly sexually active groups (HYAMS et al., 1992).

These data also indicate that false-positive anti-HCV ELISA serological results are common in the Somali population and that the second generation ELISA assay is much better than the first generation assay for identifying immunoblot positive sera. Other studies of African populations have found a high prevalence of false-positive anti-HCV ELISA results, which has been attributed to malaria or flavivirus infection, cross-reactive antibody to unknown antigens, and infection by HCV variants (ACETI

[&]quot;Western blot for human immunodeficiency virus 1.

^{&#}x27;Fluorescent treponemal antibody absorption test.

et al., 1990; Wong et al., 1990; CHAN et al., 1991; ELLIS et al., 1991; JACKSON et al., 1991; TIBBS et al., 1991;

HYAMS et al., 1993).

Our results suggest that HIV-1 infection was not common in Somalia at the time this study was conducted. A prior study in Somalia indicated that HIV-1 infection was not then widespread in that region, possibly because of minimal commerce between Somalia and the rest of Africa (SCOTT et al., 1991).

Acknowledgements

This research was supported by the US Naval Medical Research and Development Command, Bethesda, Maryland, USA, Work Units no. 3M463105H29AA335 and USA. Work Units no. 3M463105H29AA335 and 3M162787A870AR1288. The opinions and assertions herein are the private ones of the authors and are not to be construed as official or as reflecting the views of the US Navy Department, the Department of Defense, or the Somali Government.

References
Aceti, A., Taliani, G., Bac, C. & Sebastiani, A. (1990). AntiHCV false positivity in malaria. Lancet, 336, 1442-1443.
Al-Faleh, F. Z., Ayoola, E. A., Al-Jeffry, M., Al-Rashed, R.,
Al-Mofarreh, M., Arif, M., Ramia, S., Al-Karawi, M. & AlShabrawy, M. (1991). Prevalence of antibody to hepatitis C
virus among Saudi Arabian children: a community-based

virus among Saudi Arabian children: a community-based study. Hepatology, 14, 215-218.

Alter, H. J., Purcell, R. H., Shih, J. W., Melpolder, J. C., Houghton, M., Choo, Q.-L. & Kuo, G. (1989a). Detection of antibody to hepatitis C virus in prospectively followed transfusion recipients with acute and chronic non-A, non-B hepatitis. New England Journal of Medicine, 321, 1494-1500.

Alter, H. J., Coleman, P. J., Alexander, W. J., Kramer, E., Miller, J. K., Mandel, E., Hadler, S. C. & Margolis, H. S. (1989b). Importance of heterosexual activity in the transmission of hepatitis B and non-A, non-B hepatitis. Journal of the American Medical Association, 262, 1201-1295.

the American Medical Association, 262, 1201-1295.
Chan, S.-W., Simmonds, P., McOmish, F., Yap, P.-L., Mitchell, R., Dow, B. & Follett, E. (1991). Serological responses to infection with three different types of hepatitis C virus.

Lancet, 338, 1391.

Ellis, L. A., Brown, D., Conradie, J. D., Paterson, A., Sher, R., Millo, J. & Dusheiko, G. M. (1991). Regional prevalence of hepatitis C antibodies in South Africa: an analysis of fresh of nepatitis C antibodies in South Africa: an analysis of fresh and stored serum. In: Viral Hepatitis and Liver Disease, Hollinger, F. B., Lemon, S. M. & Margolis, H. S. (editors). Baltimore: Williams and Wilkins, pp. 445-447.

Esteban, J. I., Gonzalez, A., Hernandez, J. M., Viladomiu, L., Sanchez, C., Lopez-Talavera, J. C., Lucea, D., Martin-Vega, C., Vidal, W., Esteban, R. & Guardia, J. (1990). Evaluation

of antibodies to hepatitis C virus in a study of transfusion-associated hepatitis. New England Journal of Medicine, 323, 1107-1112

Everhart, J. E., Di Bisceglie, A. M., Murray, L. M., Alter, H. J., Melpolder, J. J., Kuo, G. & Hoffnagle, J. H. (1990). Risk for non-A, non-B (type C) hepatitis through sexual or household contact with chronic carriers. Annals of Internal Medicine, 112, 544-545.

Follett, E. A. C., Dow, B. C., McOmish, F., Lee Yap, P., Hughes, W., Mitchell, R. & Simmonds, P. (1991). HCV con-

firmatory testing of blood donors. Lancet, 338, 1024.
Hyams, K. C., Phillips, I. A., Moran, A. Y., Tejada, A. & Wignall, F. S. (1992). Seroprevalence of hepatitis C virus antibody in Peru. Journal of Medical Virology, 37, 127-131. Hyams, K. C., Okoth, F. A., Tukei, P. M., Vallari, D. S.,

Morrill, J. C., Long, G., Bansal, J. & Constantine, N. (1993). Inconclusive hepatitis C virus antibody results in African sera. Journal of Infectious Diseases, 167, 254-255.

Jackson, J. B., Guay, L., Goldfarb, J., Olness, K., Ndugwa, C., Mmiro, F., Kataaha, P. & Allain, J.-P. (1991). Hepatitis C virus antibody in HIV-1 infected Ugandan mothers. Lancet,

337, 551.

Saeed, A. A., Alk-Admawi, A. M., Al-Rasheed, A., Fair-clough, D., Bacchus, R., Ring, C. & Garson, J. (1991). Hepatitis C virus infection in Egyptian volunteer blood donors in

Riyadh. Lancet, 338, 459-460.

Scott, D. A., Gorwin, A. L., Constantine, N. T., Omar, M. A., Guled, A., Yusef, M., Roberts, C. R. & Watts, D. M. (1991). Low prevalence of human immunodeficiency virus-1 (HIV-1), HIV-2, and human T cell lymphotropic virus-1 infection in Somalia. American Journal of Tropical Medicine and Hygiene, 45, 653-659. Tibbs, C. J., Palmer, S. J., Coker, R., Clark, S. K., Parsons.

G. M., Hojvat, S., Peterson, D. & Banatvala, J. E. (1991). Prevalence of hepatitis C in tropical communities: the importance of confirmatory assays. Journal of Medical Virology, 34,

143-147.

Van den Hoek, J. A. R., van Haastrecht, H. J. A., Goudsmit, J., de Wolf, F. & Coutinho, R. A. (1990). Prevalence, incidence, and risk factors of hepatitis C virus infection among drug users in Amsterdam. Journal of Infectious Diseases, 162, 823-826.

Wong, D. C., Diwan, A. R., Rosen, L., Gerin, J. L., Johnson, R. G., Polito, A. & Purcell, H. (1990). Non-specificity of anti-HCV test for seroepidemiological analysis. Lancet, 336. 750-751.

Received 16 March 1993; revised 13 May 1993; accepted for publication 13 May 1993

Announcement

Second Tropical Neurology Congress Limoges, France 21-23 September 1994

The official languages of this congress will be French and English. Registration fee: FF 1500 (US \$ 250)

before 1 March 1994; FF 2000 (US \$ 325) after that date.

Further information from: Prof. M. Dumas, Institut d'Epidémiologie et de Neurologie Tropicale, Faculté de Médecine, 2 Rue du Docteur Marcland, 87025 Limoges Cedex, France. Telephone: (33) 55 43 58 20; fax: (33) 55 43 58 01.